

Containment and Prevention of MDROs in Post-acute and Long-term Care

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Speaker Disclosures

- Kara M. Jacobs Slifka, MD, MPH
 - No conflicts to disclose
 - The content of this presentation reflects my opinion and does not necessarily reflect the official position of the CDC
- Noreen Mollon, MS, CIC
 - No conflicts to disclose

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Centers for Disease Control and Prevention (CDC)





National Center for Emerging and Zoonotic Infectious Diseases ORGANIZATIONAL CHART



Division of Healthcare Quality Promotion (DHQP)

- Investigate and respond to emerging infections and adverse events in healthcare facilities
- Support the enhancement of state infrastructure for elimination of HAIs
- Develop and disseminate evidence-based guidelines and recommendations to prevent and control HAIs, antibiotic resistance, and medication errors
- Provide healthcare facilities, states, and federal agencies with data for action through the National Healthcare Safety Network (NHSN), a tool for monitoring and preventing healthcare-associated infections, used by healthcare facilities in all 50 states

Prevention & Response Branch: Long-Term Care Team

- Improve infection surveillance, prevention, and antibiotic stewardship in nursing homes
- Define and measure antibiotic use and antibiotic resistance in nursing homes
- Prevent the spread of novel and emerging resistance
- Promote NHSN reporting as a part of SNF quality measurement programs
- Provide resources and assistance to state and local health departments,
 post-acute and long-term care facilities

MDROs in Post-acute and Long-term Care (PA/LTC)

- Contain and Prevent the spread of MDROs
- Develop updated guidance specific to PA/LTC working with regulatory partners
- Provide resources and assistance to state and local health departments,
 post-acute and long-term care facilities
- Develop a better understanding of the unique challenges faced by nursing homes, especially those providing high-acuity care
- Promote the development of standardized tools and educational materials

Noreen Mollon, MS, CIC

Surveillance for Healthcare-Associated and Resistant Pathogens (SHARP) unit

Communicable Disease Division

Bureau of Epidemiology and Population Health

Michigan Department of Health and Human Services





Surveillance for Healthcare-Associated and Resistant Pathogens (SHARP) Unit

- Objectives of the SHARP Unit:
 - Coordinate activities related to Healthcare-Associated Infection (HAI) surveillance and prevention in Michigan
 - Improve surveillance and detection of antimicrobial-resistant pathogens and HAIs
 - Identify and respond to disease outbreaks
 - Use collected data to monitor trends
 - Educate healthcare providers, state and local public health partners, and the public
 - Connect partners engaged in antimicrobial stewardship activities

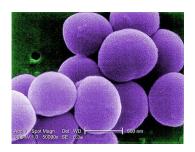


SHARP Activities

- Outbreak Response
- Infection Control Needs Assessments
- Consulting/Education
- Surveillance and Reporting
- CRE Surveillance and Prevention Initiative



Klebsiella pneumoniae



Staphylococcus aureus

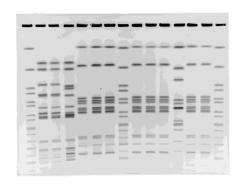


Clostridium difficile

Outbreak Response



 The MDHHS SHARP staff are available to offer our services and expertise in healthcare-associated outbreak investigations





Acinetobacter baumannii

 MDHHS can help facilities coordinate molecular testing with the MDHHS Bureau of Laboratories to identify genetic-relatedness between patient isolates (at no cost)

Session Objectives

- Discuss the public health importance of multidrug-resistant organisms (MDROs) and emerging pathogens in the postacute and long-term care settings
- Discuss risk factors for colonization and infection with MDROs
- Describe surveillance and prevention of MDROs in Michigan
- Describe strategies for preventing the spread of MDROs focused on infection prevention practices
 - Define the CDC's containment strategy
 - Discuss Infection Control Assessment and Response Tool and Michigan findings

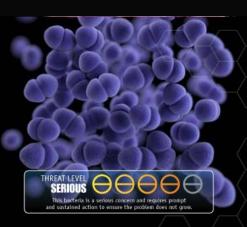
Case Example

- 70 year old admitted from a long-term acute care hospital to nursing home
 - Complicated hospital history including surgery, prolonged ICU stay, multiple courses of antibiotics
 - Spent 5 weeks in the LTACH
- On transfer, has tracheostomy, PEG tube, indwelling urinary catheter and partially healing sacral pressure ulcer
- One week later, on reviewing the chart, you find results of a culture sent from tracheostomy secretions

Case Example, continued

 Tracheostomy aspirate culture grew Klebsiella pneumoniae, >10⁵ cfu

Drug	Result		
Amikacin	Intermediate		
Ampicillin	Resistant		
Amp/Sulbactam	Resistant		
Aztreonam	Resistant		
Cefazolin	Resistant		
Cefepime	Resistant		
Ceftazidime	Resistant		
Ceftriaxone	Resistant		
Cefuroxime	Resistant		
Gentamicin	Resistant		
Levofloxcin	Resistant		
Meropenem	Resistant		
Piperacillin/Tazobactam	Resistant		
Tobramycin	Resistant		
Trimethoprim/Sulfa	Resistant		



VANCOMYCIN-RESISTANT COCCUS (VRE)



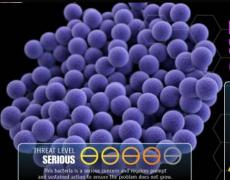


ENTEROCOCCUS INFECTIONS



SOME ENTEROCOCCUS STRAINS ARE RESISTANT TO VANCOMYCIN **LEAVING FEW OR NO TREATMENT OPTIONS**





APHYLOCOCCUS AUREUS





STAPH BACTERIA ARE A LEADING CAUSE OF

THREAT LEVEL This bacteria is a serious concern and requires prompt and sustained action to ensure the problem does not grow.







ENTEROBACTERIACEAE INFECTIONS PER YEAR







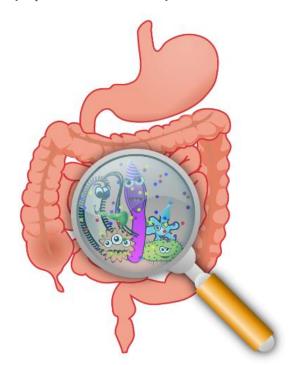


Carbapenem Resistant Enterobacteriaceae (CRE)



CRE are a public health threat

- 1. CRE cause invasive infections with high mortality (up to 40-50%)
 - Urinary Tract Infections
 - Bloodstream infections
 - Wound infections
 - Pneumonia



CRE are a public health threat

- 1. They cause invasive infections associated with high mortality rates
- Carry resistance genes on mobile genetic elements that confer high levels of resistance



Leave limited to no therapeutic options



Carbapenem-resistant Enterobacteriaceae (CRE)

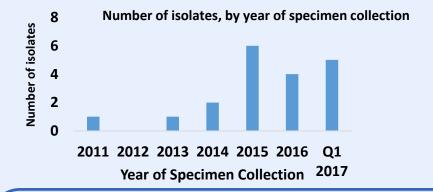
- Multiple different mechanisms can cause resistance
 - Carbapenemase-producing (CP-CRE)
 - **KPC** *Klebsiella pneumoniae* carbapenemase (most common in U.S.)
 - **NDM** New Delhi Metallo-β-lactamase
 - VIM Verona Integron-encoded Metallo- β
 - OXA Oxacillinase-48-type carbapenemase
 - **IMP** Imipenemase Metallo- β -lactamase
 - Non-carbapenemase-producing (non-CP-CRE)



Carbapenemases in other Gram negative bacteria

Proteus mirabilis, Providencia rettgeri, Citrobacter freundii

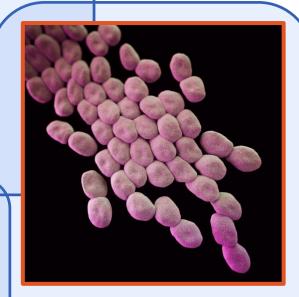
Carbapenem-Producing Organisms (CPOs)



Pseudomonas aeruginosa



VIM: 86 patients, 12 states



Acinetobacter baumannii

CPOs are a public health threat

- 1. They cause invasive infections associated with high mortality rates
- 2. Carry resistance genes on mobile genetic elements that confer high levels of resistance
- 3. CRE have spread throughout the United states and other countries and have the potential to spread more widely

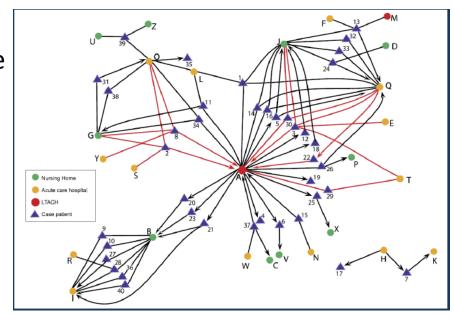


PROBLEM:

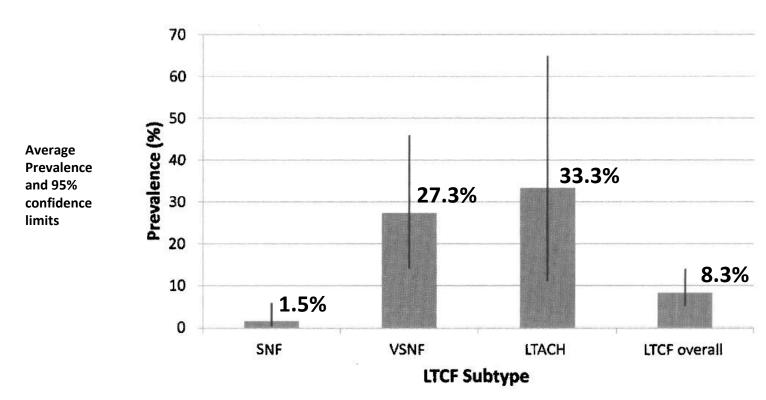
Antibiotic-resistant germs can spread like wildfire.

Healthcare networks driving outbreaks: Findings from public health investigations

- Post-acute care facilities with longer length of stay and high acuity of care (e.g., ventilator services, IV therapy, wound care) expand the burden of resistance within a region
- Gaps in IPC program infrastructure and practices can augment this problem



Carriage of CP-CRE (*Klebsiella* pneumoniae) among Hospitalized patients admitted from Post-acute/Long-term care, 2012



Older adults are at high risk for infections with MDROs



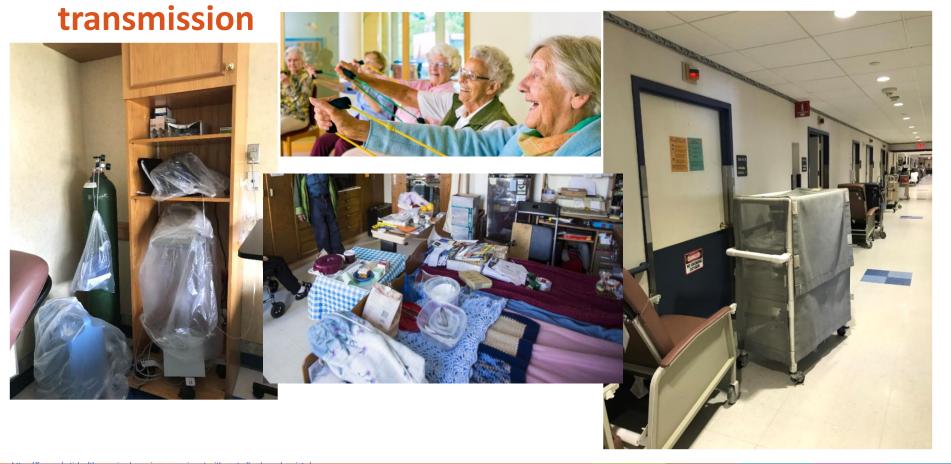


Risk Factors for colonization with MDROs

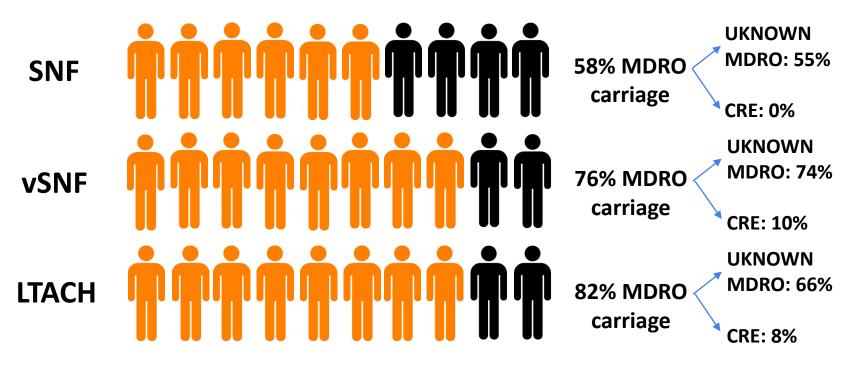
- Indwelling medical device (urinary catheter, PEG tube, trach, central line)
- Lower functional status
- Presence of wounds or decubitus ulcers

- Antibiotic use in prior 3 months
- Fluoroquinolone use
- History of hospitalization
- Older age
- Comorbid medical conditions

Nursing home setting provides opportunity for



Carriage of ANY MDRO (Median %)

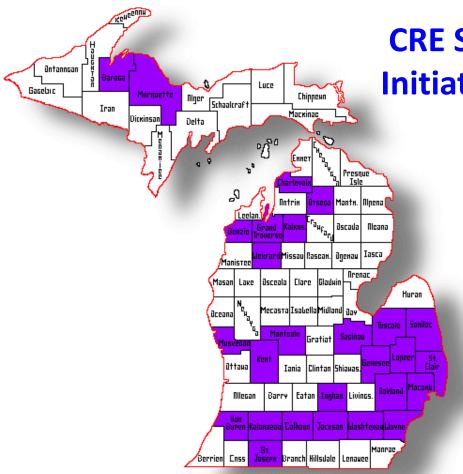


Carbapenem-resistant *Enterobacteriaceae*Surveillance and Prevention Initiative

- Began in 2012
- Voluntary reporting of CRE
 - Klebsiella pneumoniae and Escherichia coli resistant to any carbapenem (Sept 2012-Aug 2017)
 - Klebsiella spp., Enterobacter spp., Escherichia coli positive for carbapenemase production by a phenotypic or molecular test or those resistant to ANY carbapenem if no confirmatory testing done (Sept 2017 – current)
- Implementation of a CRE prevention plan
 - Facility-specific based on needs and resources
 - Examples: policy/procedure changes, education, communication, compliance monitoring (hand hygiene, contact precautions), CHG bathing

CRE Surveillance and Prevention Initiative Voluntary Participation

	Baseline Period	Intervention Period	Acute Care	LTAC	LTC/SNF	Total
Phase 1	Sept 2012-Feb 2013	Mar 2013- Aug 2014	17	4	0	21
Phase 2	Mar 2014-Aug 2014	Sept 2014-Feb 2016	7	2	0	9
Phase 3	Sept 2015-Feb 2016	Mar 2016-Aug 2017	4	4	2	10
New facilities	Sept 2017-Feb 2018	Mar 2018-Aug 2019	14	7	0	21
Combined Cohort	Sept 2017-Feb 2018	Mar 2018-Aug 2019	42	17	2	61

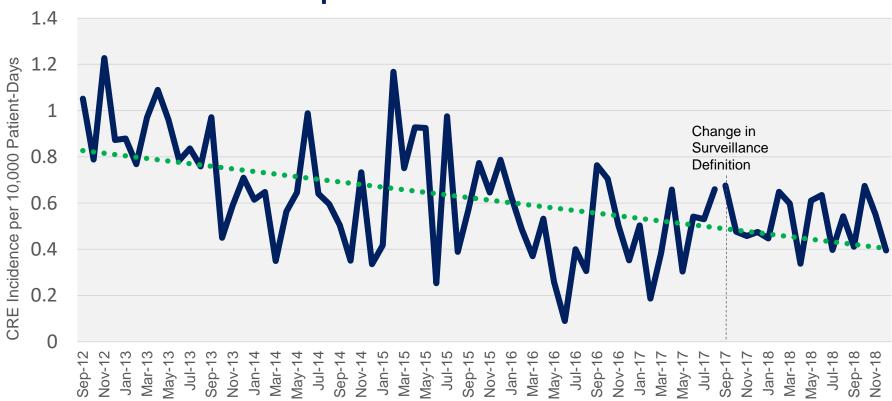


CRE Surveillance and Prevention Initiative Participation by County, Sept 2018

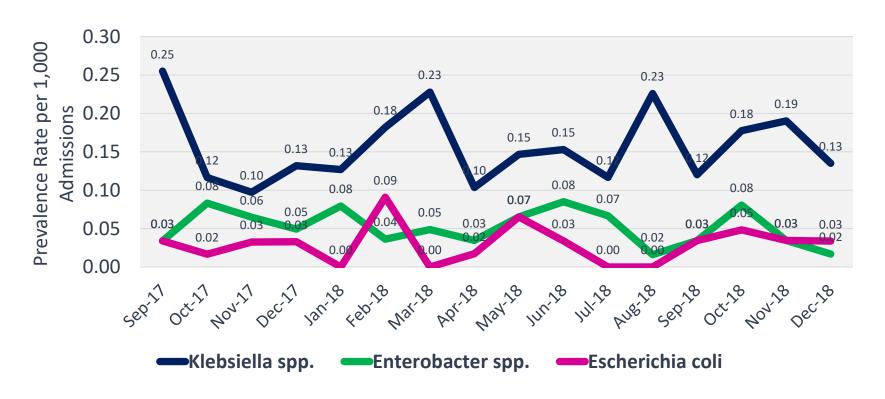
61 Facilities in 27 counties

Overall Inpatient CRE Incidence

Sept 2012 - Dec 2018



Inpatient CRE Prevalence Rate by Organism Sept 2017 – Dec 2018



Carbapenemase-producing CRE Reporting

- Reportable disease in Michigan starting January 2018
- Surveillance definition endorsed by CSTE/CDC
- CP-CRE cases are reported using the Michigan Disease Surveillance System (MDSS)
 - Web-based communicable disease reporting system for the state of Michigan
 - Cases can be reported by:
 - Electronic laboratory report (ELR)
 - Manual case entry

CP-CRE Reporting Requirements

- Laboratories, infection prevention and Local Health Departments are required to report all cases of CP-CRE according to the following criterion for Klebsiella spp., E. coli, or Enterobacter spp.:
 - Healthcare record contains a diagnosis of Carbapenemase-producing Carbapenem-resistant Enterobacteriaceae (CP-CRE), KPC, NDM, OXA-48, IMP or VIM or other novel carbapenemase
 - Any isolate of *Klebsiella* spp., *E. coli*, or *Enterobacter* spp. demonstrating carbapenemase production by a phenotypic test (e.g., Carba NP, CIM, mCIM)
 - Any isolate of Klebsiella spp., E. coli, or Enterobacter spp. with a known carbapenemase resistance mechanism (e.g., KPC, NDM, OXA-48, IMP, VIM, or other carbapenemase gene) by a recognized molecular test (e.g., PCR, Expert Carba-R)

CP-CRE Reporting Requirements

- If laboratories are <u>unable to detect</u> **CP-CRE**, (*i.e.*, cannot test for carbapenemase production (phenotypic) or resistance mechanism (molecular test):
 - Report any isolate of Klebsiella spp., E. coli, or Enterobacter spp. with a minimum inhibitory concentration (MIC) of any of the following:
 - ≥4 mcg/ml for Meropenem
 - ≥4 mcg/ml Imipenem
 - ≥4 mcg/ml Doripenem
 - ≥ 2 mcg/ml for Ertapenem

Case Classification

CONFIRMED CP-CRE

- Klebsiella spp., E. coli, Enterobacter spp.
 - Positive phenotypic test OR
 - Positive carbapenem resistance mechanism

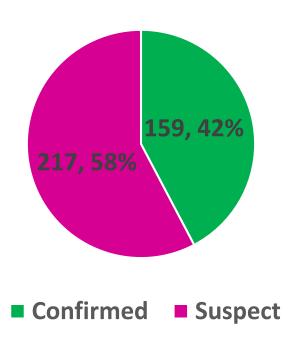
SUSPECT CP-CRE

- Klebsiella spp., E. coli, Enterobacter spp.
 - Resistance to at least 1 carbapenem
 - No phenotypic or molecular testing done

NOT a CASE

- BOL report is negative for phenotypic and molecular tests
- All carbapenems are susceptible (MICs don't match case definition)
- Not Enterobactericeae

CP-CRE Cases Reported to MDSSJan – Dec 2018



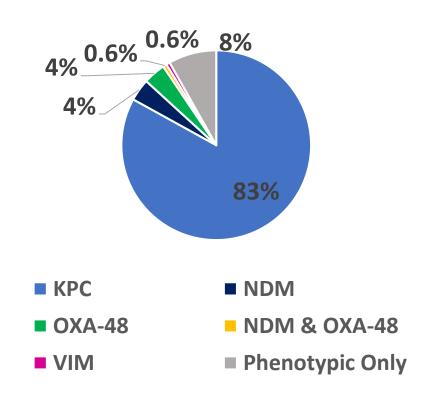
CP-CRE Cases by Organism

Jan - Dec 2018

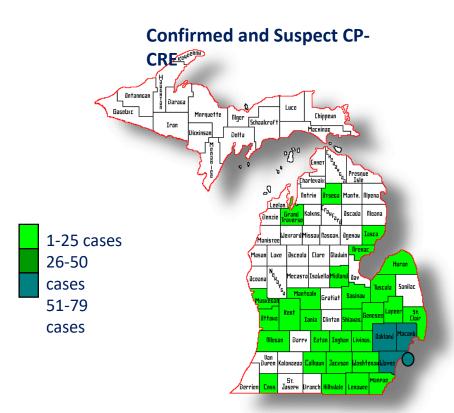
	CP-CRE Cases		
	Confirmed	Suspect	Total
Organism	n=159	n=217	n=376
Klebsiella spp.	110 (69%)	89 (41%)	199 (53%)
Klebsiella pneumoniae	102	68	170
Klebsiella aerogenes	4	14	18
Klebsiella oxytoca	3	7	10
Klebsiella variicola	1	0	1
Escherichia coli	23 (14%)	69 (32%)	92 (42%)
Enterobacter spp.	26 (16%)	36 (17%)	85 (23%)
Enterobacter cloacae	26	57	83
Enterobacter asburiae	0	1	1
Enterobacter hormaechei	0	1	1

Confirmed CP-CREJan – Dec 2018

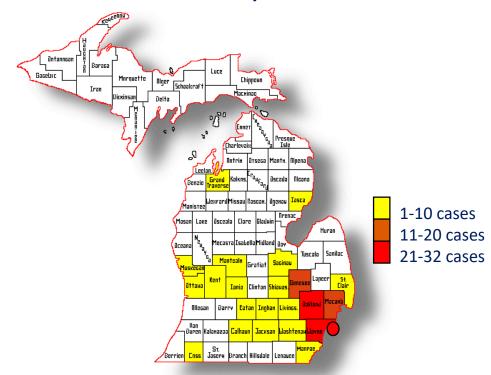
- 159 positive for carbapenemase production by a phenotypic test (e.g., mCIM, MHT)
- 146 results available for resistance mechanism
 - 132 KPC
 - 6 NDM-1
 - 6 OXA-48
 - 1 NDM-1 & OXA-48
 - 1 VIM



CP-CRE Cases by County*, 2018



Confirmed CP-CRE only



*based on county of residence

CP-CRE and **Novel** Resistance Activity

Carbapenemases:

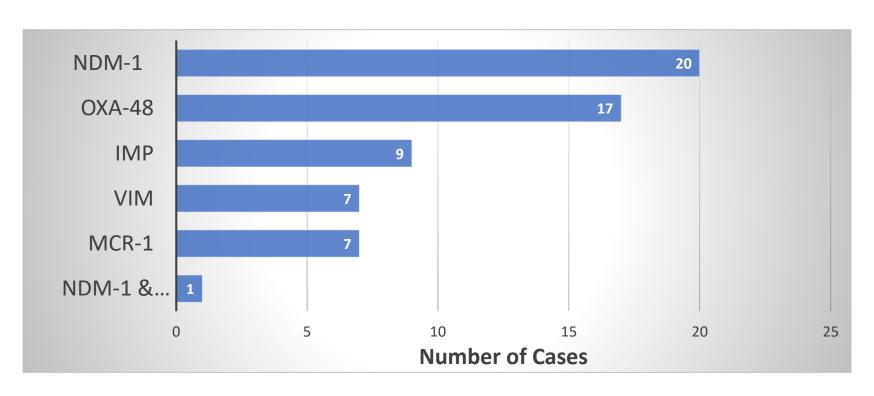
- Klebsiella pneumoniae carbapenemase (KPC)
- New Delhi metallo-β-lactamase (NDM)
- Verona integron encoded metallo-β-lactamase (VIM)
- Imipenemase metallo-β-lactamase (IMP)
- Oxacillinase-48 (OXA-48)

Other resistance elements:

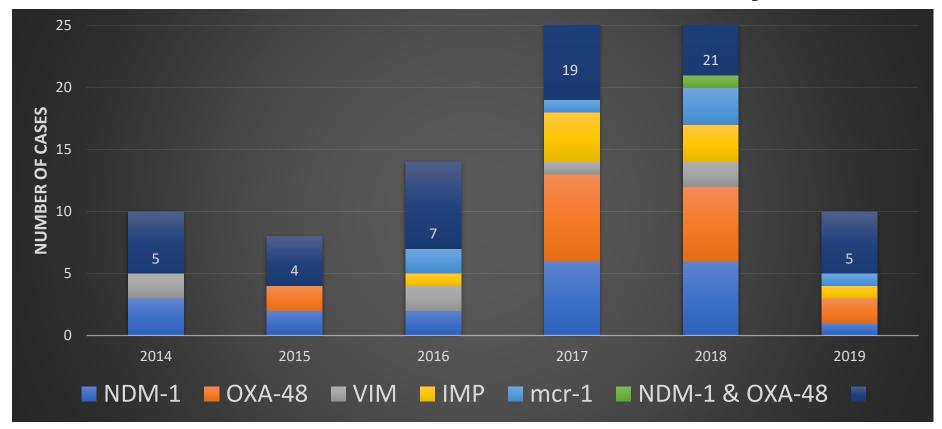
Mobile colistin resistance (mcr)

Confirmed Novel Resistance Cases

2014 - Current



Confirmed Novel Resistance Cases by Year



Novel Resistance Case Demographics







Median Age, 62 yrs

Male, 32 (52%)

Comorbid conditions

Cardiovascular disease – 39%

Diabetes – 33%

Chronic lung disease – 21%

Renal failure, chronic wound – 20%

Malignancy, vent dependent -11%

Urinary catheter – 10%

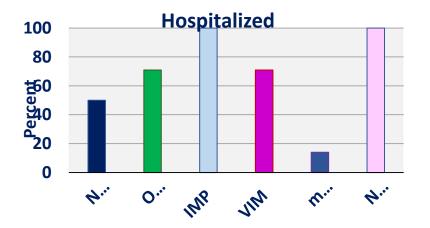
dementia - 7%

(range 5 - 87 yrs)

Common Risk Factors for Novel Resistance

Healthcare Exposures

 62% were hospitalized 6 months prior to positive culture



International Travel

 51% had travel 6 months prior to positive culture



Carbapenemase and Resistance Mechanism Testing

- Laboratories are strongly encouraged to submit CRE isolates to the MDHHS Bureau of Laboratories
 - Confirm organism identification
 - Perform modified carbapenem inactivation method (mCIM) testing
 - Perform PCR testing for KPC, NDM, OXA-48 like, IMP, VIM
 - If mCIM or PCR are positive, antimicrobial susceptibility testing (AST) will be performed





THERE'S A FUNGUS

AMONG US.

App labilities 20

Candida can cause serious infections

- Candidemia is the most common HAI bloodstream infection.
- 30% mortality
- Risk factors include:
 - Broad-spectrum antibiotic use
 - Central venous catheters
 - Immune compromise



1. Often misidentified



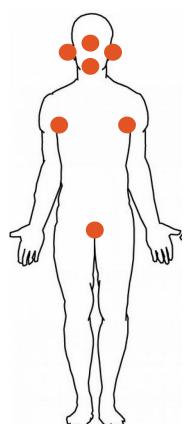
- Often misidentified
- 2. Resistant to antifungal drugs



- 1. Often misidentified
- 2. Resistant to antifungal drugs
- 3. Causes invasive infections with high mortality



C. auris Colonizes Skin and Other Body Sites



Colonization poses a risk for:

- Invasive infection
- Transmission to others

Risk Factors for Candida auris

- Older age
- Multiple healthcare stays (post-acute and long term)
- Prolonged healthcare stay
- Taking antibiotics and antifungals
- Tracheostomy
- Ventilator
- Feeding tubes
- Central lines



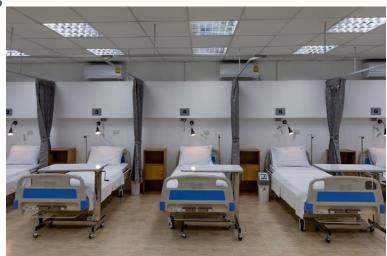
Candida auris colonizes the environment



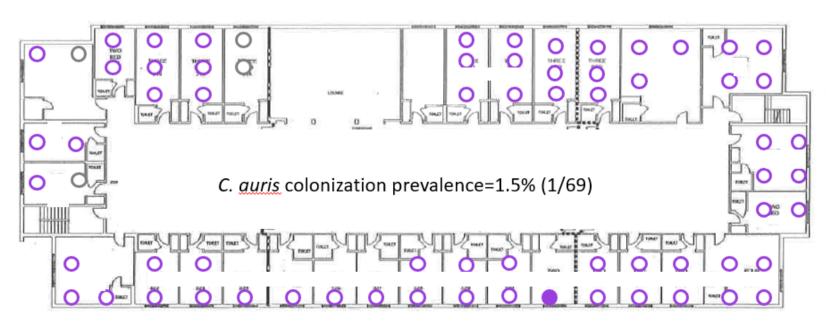
https://5.imimg.com

- 1. Often misidentified
- 2. Resistant to antifungal drugs
- 3. Causes invasive infections with high mortality
- 4. Can cause outbreaks in healthcare settings,

All the makings of a fungal superbug!

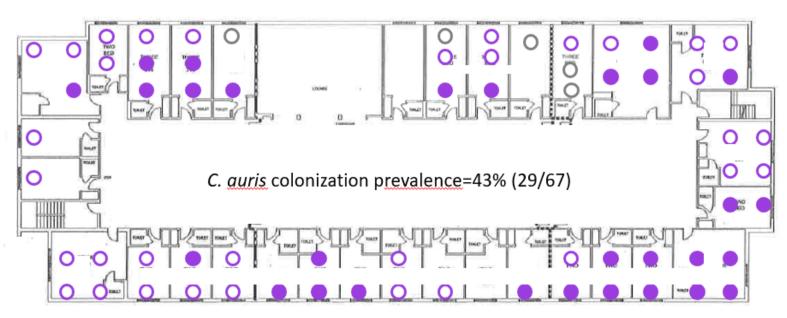


vSNF A Ventilator/Trach Floor March 2017 *C. auris* PPS Results



- C. auris positive
- Screened negative for C. auris
- Not tested for C. <u>auris</u> (refused or not in room)

vSNF A Ventilator/Trach Floor January 2018 *C. auris* PPS Results

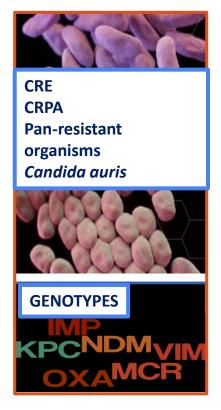


- C. auris positive
- Screened negative for C. auris
- O Not tested for C. auris (refused or not in room)

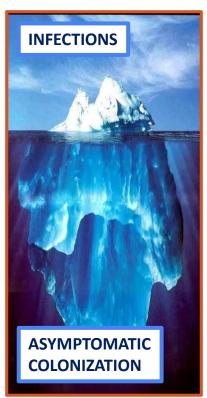
Slide courtesy of Chicago Department of Public Health.

Characteristics of MDROs in PA/LTC

Resistance



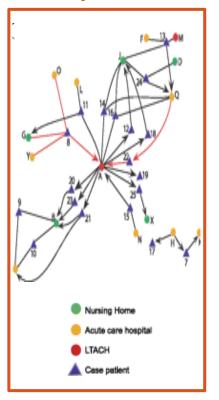
Detection



Transmission

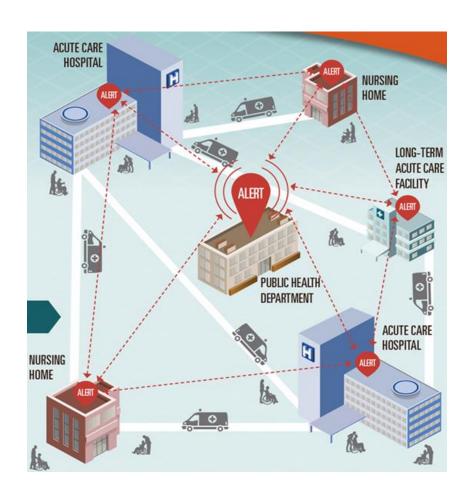


Spread



Candida auris in Michigan

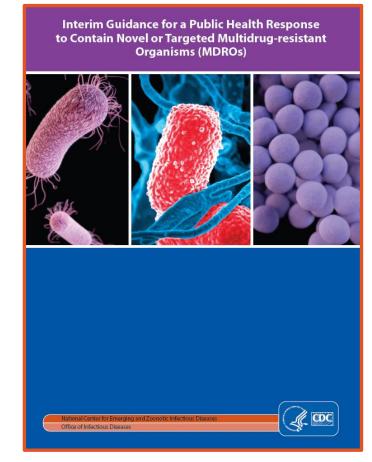
- In 2019, CSTE and CDC passed a position statement to make *C. auris* nationally notifiable. Michigan followed suit and as of January 1, 2019 it is reportable in Michigan
- Please report any patient or laboratory finding to MDHHS that meets either of the following criteria:
 - Detection of *C. auris* in a specimen using either culture or a culture independent diagnostic test (CIDT) (e.g., Polymerase Chain Reaction [PCR])
 - Detection of an organism that commonly represents at a *C. auris* misidentification in a specimen by culture (i.e., *Candida haemulonii*): https://www.cdc.gov/fungal/diseases/candidiasis/pdf/Testing-algorithm-by-Method-temp.pdf
- The important thing to note is Candida auris is bad. This is not your average yeast. This will require extensive investigation. https://www.cdc.gov/fungal/candida-auris/tracking-c-auris.html





Containment and Prevention of MDROs

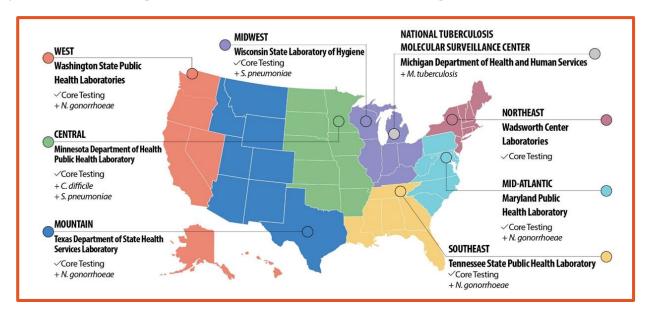
- Systematic approach to slow spread of novel or rare multidrug-resistant organisms or mechanisms through aggressive response to ≥1 case
 - Pan-resistant organisms
 - Carbapenemase-producing organisms
 - mcr-1
 - Candida auris
- Response based on pathogen/resistance mechanism



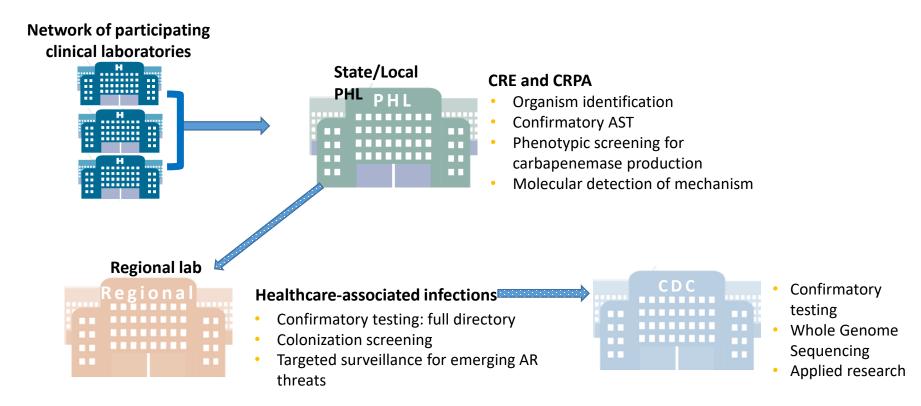
https://www.cdc.gov/hai/outbreaks/mdro/index.html

Antibiotic Resistance Laboratory Network (ARLN)

- Tiered network established in 2016 to support nationwide lab capacity to rapidly detect antibiotic resistance in healthcare, food, and the community
- Public health laboratories in 50 states, 6 cities and Puerto Rico
 - Carbapenemase testing for CRE and CR-Pseudomonas aeruginosa



ARLN: Enhanced Capacity Through Tiered Testing



MDHHS Bureau of Labs



- Bureau of Laboratories has expanded test offerings to include:
 - Enterobacteriaceae, Acinetobacter, and Pseudomonas aeruginosa
 - Confirmation of carbapenemase production and colistin resistance
 - Genetic markers for KPC, NDM, VIM, OXA-48, and MCR-1
 - Perform modified carbapenem inactivation method (mCIM) testing

Systematic public health response to slow the spread of emerging AR



Systematic public health response to slow the spread of emerging AR



Single case of emerging resistance

Systematic public health response to slow the spread of emerging AR



Onsite assessment using standardized tools

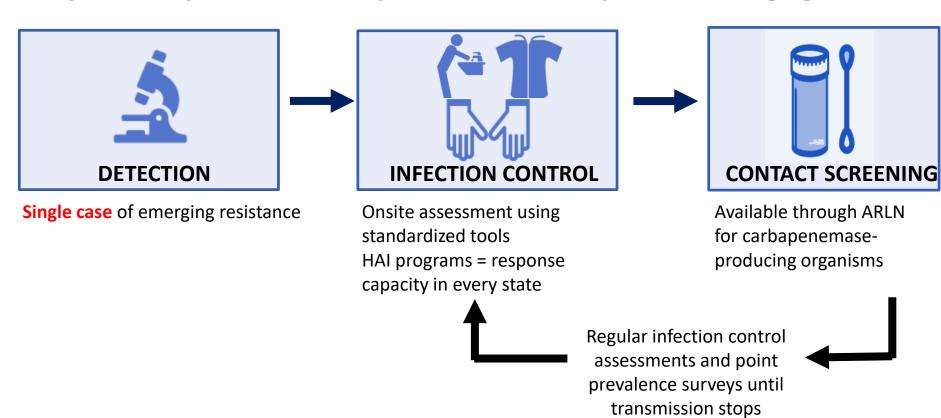
HAI programs = response capacity in every state

Systematic public health response to slow the spread of emerging AR



Available through ARLN for carbapenemase-producing organisms

Systematic public health response to slow the spread of emerging AR



What to expect during a response?

- You have a critical role in containing emerging antibiotic resistance
- If unusual resistance identified in a resident at your facility or who has been in your facility
 - The health department will reach out about infection control assessments (ICAR) and contact screening
 - Focus is on preventing spread of resistance



Common Themes from CRE and CRPA Responses

- Residents in long length of stay, high acuity settings at highest risk
- Factors in transmission
 - Gaps in adherence to hand hygiene and Contact Precautions
 - Environmental contamination, including improperly cleaned equipment from contracted providers
 - Resident supplies in sink splash zone
 - Failure to communicate resident status at transfer
- Larger clusters take longer to control
 - Multiple on-site visits to observe infection control and multiple rounds of PPS
 - Staff training on hand hygiene, PPE use, environmental cleaning

ICAR Goals

- Increase patient safety
- Expand infection control resources
- Increase the number of infection control consultations provided by the SHARP unit



Methods

Used a CDC tool to conduct infection control needs assessments

- Review facility practices:
 - Infection Control Infrastructure
 - Infection Control Training, Competency, and Implementation of Policies and Practices
 - Systems to Detect, Prevent and Respond to Healthcare-Associated Infections and Multi-Drug Resistant Organisms

The CDC Evaluation Tool

Organized into 4 sections:

- 1. Facility demographics
- 2. Infection control program and infrastructure
 - 9 domains
- 3. Direct observation of facility practices (optional)
- 4. Infection control guidelines and other resources



Assessment and Response



Discuss findings with Infection
Preventionists and other staff



Report facility findings back to facility leadership



Aggregate findings

Strengths

Areas for opportunity

Facility Recruitment: 2015-2018

- Voluntary participation
- Collaborative, NOT regulatory
- Advertised to interested facilities:
 - Website, flyers, emails
 - Professional societies (e.g. MSIPC, APIC GL, HCAM)
 - Meetings and conference presentations



Facility recruitment: 2019-

- Response to HAI outbreak
- Response to identification of a novel organism
- Volunteer!

Participating LTC Facilities

- 41 assessments completed in LTC from 2015-2018
- 28 (68%) assessments completed on-site
- All facilities were licensed by the state
- 39 (95%) were certified by CMS
- Mean licensed beds: 110 beds (range 46-260)
- Staff hours per week dedicated to IP: 22.4 hours (range 2-40)

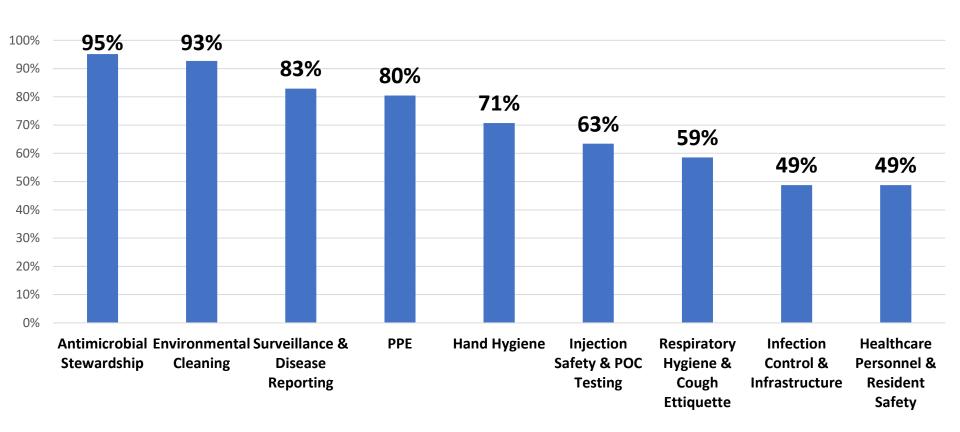


ICAR Results



- Gaps were common
- Assessments identified at least 1 gap in each facility

Facilities with at Least 1 Gap by Domain



Recommendations

- Antimicrobial Stewardship
 - 30 (73%) Develop policies and procedures
 - Develop education and training for staff



- Environmental Cleaning
 - 32 (78%) Develop policies and procedures for cleaning
 - 17 (41%) Improve regular training programs to include all staff that clean
 - 17 (41%) Develop audit/feedback process for cleaning
- Surveillance & Disease Reporting
 - 21 (51%) Develop policies and procedures for conducting surveillance

Recommendations

- PPE
 - 28 (68%) Develop an audit/feedback process not just for contact precautions
- Hand Hygiene
 - 21 (51%) Provide feedback from audits, facility-level and individual-level
 - 15 (37%) Start more formal audit program
- Injection Safety & Point of Care Testing
 - 21 (51%) Develop a formal audit program
 - 20 (49%) Develop a formal feedback program
 - 16 (39%) Implement competency-based trainings



Recommendations

- Respiratory hygiene/cough etiquette
 - 21 (51%) Implement Health Department recommendations for signage

- Infection Control Program & Infrastructure
 - Specific training in infection control for IP staff

- Healthcare personnel & Resident Safety
 - 13 (32%) Develop or update policies and procedures for TB testing/screening, HCW influenza vaccination



Lessons Learned



No program is perfect—always room for improvement



Infection
prevention involves
a lot of
departments- get
to know your
colleagues!



ICAR is a great tool and free resource to enhance your program

How Can ICAR Help You?

- ✓ Collaborative process, NOT regulatory
- √ Focus on quality improvement
- √ Free consultation
- ✓ Strengthen your IP program
- ✓ Add another tool to your toolbox





Facility Level Prevention Strategies



Hand hygiene



Personal
Protective
Equipment and
Precautions



Meticulous environmental disinfection



Hand Hygiene



Barriers to Hand Hygiene (HH) adherence in NHs

Workload

Access

Guidelines

Confusion with gloves

Lack of Education

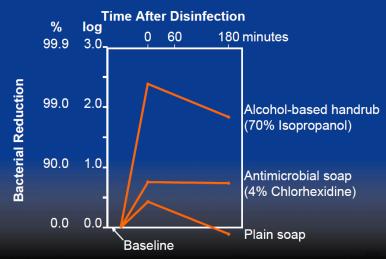
Barriers to Hand Hygiene (HH) adherence in NHs

Workload Forgot HH because of workload 27.5% lack of alcohol-based hand Access rub Belief that HH guidelines **Guidelines** aren't applicable in LTC **Confusion with gloves** No HH because of glove use 55% never/rarely received **Lack of Education** personal feedback on HH practices Ashraf MS et al. ICHE 2010; 31(7):758-762

Efficacy of Hand Hygiene Preparations in Killing Bacteria



Ability of Hand Hygiene Agents to Reduce Bacteria on Hands



Adapted from: Hosp Epidemiol Infect Control, 2nd Edition, 1999.

The Truth about HH



If hands are not visibly soiled, use an alcohol-based hand rub (ABHR)





§483.80 Infection Control

Literature:

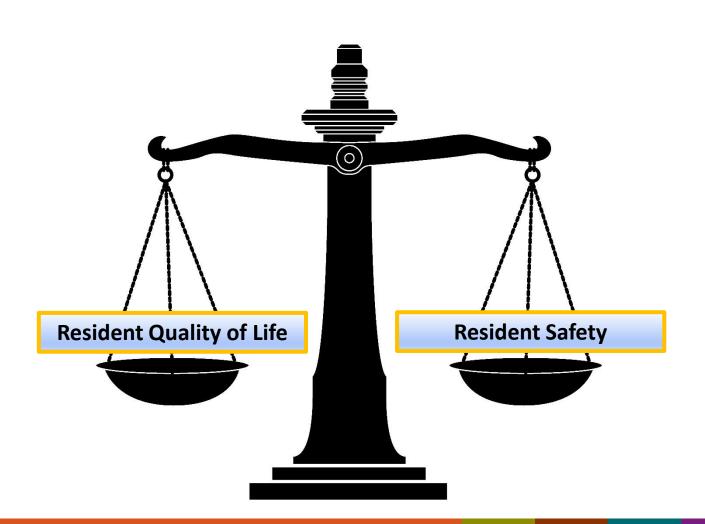
ABHR is a faster, more convenient, less drying method of HH for HCWs in a LTCF AND it improved compliance. ABHR was more efficacious than soap and water in removing pathogens already present on HCW hands.

Mody L. et al. ICHE 2003; 24(3):165-171

"consistent with accepted standards of practice such as the use of ABHR instead of soap and water in all clinical situations except when hands are visibly soiled (e.g., blood, body fluids), or after caring for a resident with known or suspected C. difficile or norovirus infection during an outbreak, or if infection rates of C. difficile are high..."



Personal Protective Equipment & Precautions





BREAK THE CHAIN!

- √ Immunizations
- Treatment of underlying disease
- Health insurance
- Patient education



BREAK THE CHAIN!

- ✓ Diagnosis and treatment
- Antimicrobial stewardship

BREAK THE CHAIN!

- Cleaning, disinfection, sterilization
- Infection prevention policies
- ✓ Pest control



Infectious agent

· Bacteria · Fungi · Viruses · Parasites



Susceptible host

· Any person, especially those receiving healthcare.

Reservoir

- Dirty surfaces Animals/ and equipment insects
- · People · Soil (earth)







Portal of entry

- . Broken skin/incisions Respiratory tract
- Mucous membranes
- . Catheters and tubes

Portal of exit

- . Open wounds/skin





Mode of transmission

· Contact · Ingestion (direct or . Inhalation indirect)



BREAK THE CHAIN!

- Hand hygiene
- ✓ Personal protective equipment
- Personal hygiene
- ✓ First aid
- / Removal of catheters and tubes

BREAK THE CHAIN!

- Hand hygiene
- √ Personal protective equipment
- Food safety
- Cleaning, disinfection, sterilization
- √ Isolation

BREAK THE CHAIN!

- Hand hygiene
- Personal protective equipment
- ✓ Control of aerosols and splatter
- Respiratory etiquette
- Waste disposal



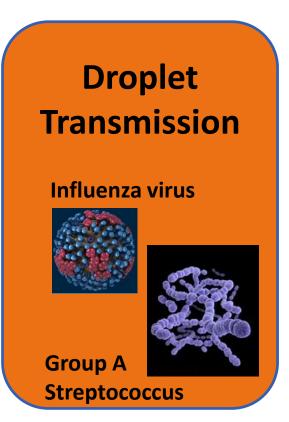


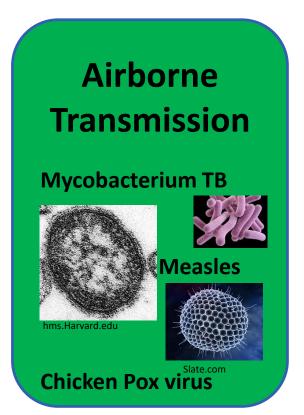
Routes of Transmission of Infectious Pathogens

Direct and Indirect Contact
Transmission

Hepatitis B

C.diff





Standard Precautions

Group of infection prevention practices

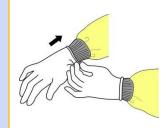
Hand Hygiene	Respiratory hygiene and cough etiquette	
Personal Protective Equipment	nent Environmental cleaning and disinfection Reprocessing of reusable medical equipment	
Safe injection practices		

- Applies to all residents regardless of suspected or confirmed infection status
- All blood, body fluids, secretions, excretions except sweat, non-intact skin, and mucous membranes may contain transmissible infectious agents



Standard & Transmission-Based Precautions

7	Standard Precautions	 Hand hygiene PPE Safe injection practices Respiratory hygiene and cough etiquette Environmental cleaning and disinfection Reprocessing of reusable medical equipment
٥	Transmission-Based Precautions	Contact PrecautionsDroplet Precautions





Airborne Precautions

Transmission-Based Precautions

Contact Precautions





Transmission-Based Precautions

- Perform hand hygiene
- PPE donned before room entry
- PPE doffed and hand hygiene performed before room exit or provided care for another resident

- Ideally resident placed in private room
- Consider cohorting
- Clear signage, easy access to ABHR, PPE, restock supplies





Standard Precautions

Hand hygiene

- PPE
- Safe injection practices
- Respiratory hygiene and cough etiquette
- Environmental cleaning and disinfection
 - Reprocessing of reusable medical equipment

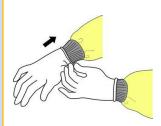


- Droplet Precautions
- Airborne Precautions



Transmission-Based Precautions





CDC 2007 Guideline for Isolation Precautions: https://www.cdc.gov/hai/pdfs/isolation2007.pdf HICPAC Core Infection Prevention and Control Practices for Safe Healthcare Delivery in All Settings: https://www.cdc.gov/hicpac/recommendations/core-practices.html

Enhanced Barrier Precautions

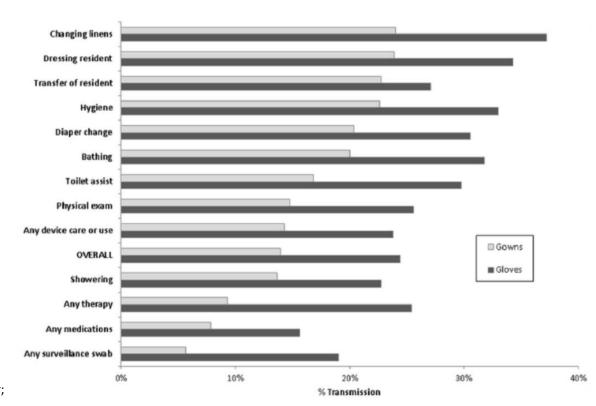
- The use gowns and gloves during high-contact resident care activities
 - Dressing
 - Bathing
 - Transferring
 - Providing hygiene
 - Changing linens
 - Changing briefs or assisting with toileting
 - Device care or use of a device (urinary catheter, central line, feeding tube, tracheostomy)
 - Wound care (any skin opening requiring a dressing)



Mary-Claire Roghmann

MRSA Transmission to Gowns and Gloves of HCW during care of MRSA colonized residents

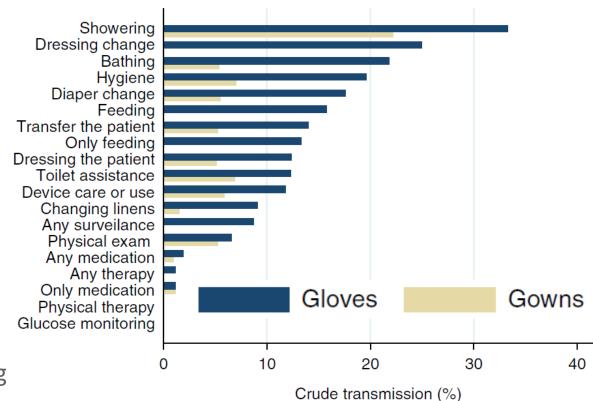
- Highest Risk:
 - Dressing
 - Transferring
 - Providing hygiene
 - Changing linens
 - Toileting
- Lowest Risk:
 - Giving meds
 - Glucose monitoring



Roghmann et al. Infect Control Hosp Epidemiol. 2015 September; 36(9): 1050-1057

Resistant Gram-negative Bacteria (RGNB) Transmission to Gowns and Gloves of HCW during care of RGNB colonized residents

- Highest Risk:
 - Showering
 - Hygiene
 - Toileting
 - Wound dressing changes
- Lowest Risk:
 - Assist feeding
 - Giving meds
 - Glucose monitoring



When to use Contact Precautions for MDRO colonized or infected residents

 Wounds, secretions, or excretions that are unable to be covered or contained,

- For preventing spread of rare and highly resistant pathogens,
- On units or in facilities where, despite attempts to control the spread of MDROs, ongoing transmission is documented or suspected.

	Applies to:	PPE used for these situations:	Required PPE	Room restriction
Contact Precautions	All residents infected or colonized with MDROs in specific situations	 Presence of wounds, secretions, or excretions that are unable to be covered or contained With rare and highly resistant pathogens (novel, pan-resistant) On units or in facilities where, despite attempts to control the spread of MDROs, ongoing transmission is documented or suspected 	Gloves and gown on EVERY room entry	Yes, except for medically necessary care.
Enhanced Barrier Precautions	All residents infected or colonized with MDROs when Contact Precautions does not apply	During high-contact resident care activities: Dressing Bathing Transferring Providing hygiene Changing linens Changing briefs or assisting with toileting Device care or use of a device: central line, urinary catheter, feeding tube, tracheostomy Wound care: any skin opening requiring a dressing	Gloves and gown (must change between residents)	None.



Cleaning & Disinfection

Room for Improvement: Environmental Cleaning

- Multiple use devices reused without cleaning
- Insufficient time for cleaning/disinfection given staffing constraints
- Proximity of resident supplies to sink and toilet
- Inappropriately performed terminal cleaning
- Insufficient contact time after using wipes
- Lapses regarding separation of clean/dirty



Case Example

- 79 year old resident is admitted to an acute care hospital from the nursing home with a urinary tract infection
- Short-stay resident on nursing home's skilled nursing unit for wound care
- Medical History: Type 2 Diabetes mellitus, hypertension, left leg wound, urinary retention requiring urinary catheter
- Urine culture on admission grows Acinetobacter resistant to Carbapenem antibiotics
- Further testing indicates OXA-23 carbapenemase production

Case Example, continued

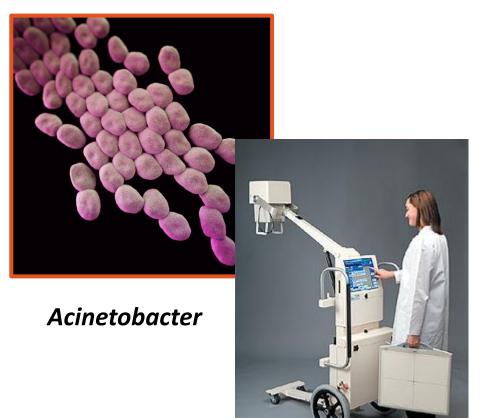
- Health department notifies nursing home of laboratory result and recommends an investigation
 - Resident had no prior MDROs; not in Contact Precautions, has roommate
 - Laboratory lookback: 2 reports of resistant Acinetobacter
 - Point Prevalence survey: 3 residents with OXA-23
 - ICAR

IX.	Environmental Cleaning	
	Assessment	
A.	The facility has written cleaning/disinfection policies which	
	include routine and terminal cleaning and disinfection of	O Yes O

Case Example: ICAR Results

- Trained, experienced IP
- ABHR and gloves available immediately inside of every resident room
- Early stages of starting an auditing & feedback program for hand hygiene and environmental services

- Limited access to gowns
- Confusion over responsibility for cleaning shared equipment
- Limited access to cleaning & disinfectant wipes



Kwipped.com



Transitions of Care

		NFECTION CON				
S		Patient/Resident		nation to resolving re	Discharg	
Demographic	Sending Facility Name:		Contact Name	:	Contact Phone:	
Dem	Receiving Facility Name:					
Precautions	Currently in Isolation Precautions? Yes If Yes check: Contact Droplet Airborne Other:				No Isolation Precautions	
	Did or does have (send o	locumentation):		Current Infection		
	Multiple Drug Resistant Org	ganism (MDRO):		☐ Yes		
	MRSA					

Facility-level Prevention

- Surveillance: Be aware of MDROs
- Policies and procedures: infection prevention, EVS, Resident & staff health programs
- Education & competency-based training for healthcare providers
- Communication at transitions of care
- Minimize use of invasive devices, appropriate device care
- Promote antibiotic stewardship

- Use your resources!
- Engagement at all levels is essential

CDC Nursing Home IP Training Course

- 23 web-based, self-study modules; close to 20 CE hours
- Curriculum designed to cover the core activities and practices of a NH IPC program
- Based on CDC guidance and best-practice recommendations
- Target audience nursing home staff given responsibility for IPC program implementation

https://www.cms.gov/Medicare/Provider-Enrollment-and-Certification/SurveyCertificationGenInfo/Downloads/QSO19-10-NH.pdf

DEPARTMENT OF HEALTH & HUMAN SERVICES
Centers for Medicare & Medicaid Services
7500 Security Boulevard, Mail Stop C2-21-16
Baltimore, Maryland 21244-1850



Center for Clinical Standards and Quality/Quality, Safety & Oversight Group

Ref: QSO-19-10-NH

DATE: March 11, 2019

TO: State Survey Agency Directors

FROM: Director

Quality, Safety & Oversight Group

SUBJECT: Specialized Infection Prevention and Control Training for Nursing Home Staff in

the Long-Term Care Setting is Now Available

Memorandum Summary

- The Centers for Medicare & Medicaid Services (CMS) and the Centers for Disease Control and Prevention (CDC) collaborated on the development of a free on-line training course in infection prevention and control for nursing home staff in the long-term care setting.
- The training provides approximately 19 hours of continuing education credits as well as a certificate of completion.
 - The "Nursing Home Infection Preventionist Training Course" is located on CDC's TRAIN website (https://www.train.org/cdctrain/training_plan/3814).
- This mame supersades mame Quality Sefety & Oversight policy memorandum OSO

Lessons Learned and Moving Forward

- How can we better address prevention and containment of MDROs?
- What steps have you taken?
- Roadblocks? Successes?
- How can we provide further support?
- What resources would be most useful?
- Feedback on Enhanced Barrier Precautions

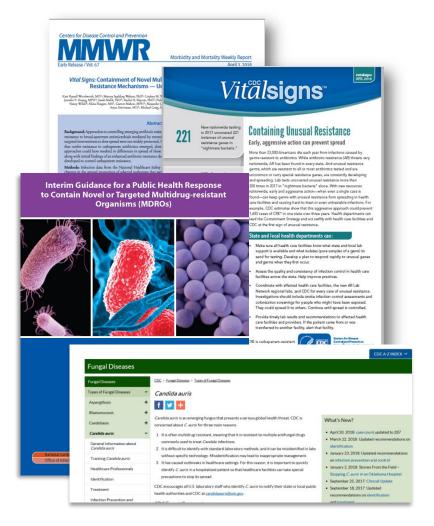


What Facilities Can Do

- Plan for unusual resistance arriving in your facility. Find resources: www.cdc.gov/hai/outbreaks/mdro
- Leadership: Work with the health department to stop spread of unusual resistance.
 Review and support infection control in the facility.
- Clinical labs: Know what isolates to send for testing. Establish protocols that immediately notify the health department, healthcare provider, and infection control staff of unusual resistance. Validate new tests to identify the latest threats. If needed, use isolates from www.cdc.gov/ARIsolateBank.
- Healthcare providers, epidemiologists, and infection control staff: Place patients with unusual resistance on contact precautions, assess and enhance infection control, and work with the health department to screen others. Communicate about status when patients are transferred. Continue infection control assessments and colonization screenings until spread is controlled. Ask about any recent travel or health care to identify at-risk patients.

Resources

- Interim Guidance to Contain Novel MDROs
 - https://www.cdc.gov/hai/containment/guidelines.html
- CDC CRE Toolkit
 - https://www.cdc.gov/hai/containment/guidelines.html
- Vital Signs on Containment
 - https://www.cdc.gov/mmwr/volumes/67/wr/mm6713e1.htm?s_cid=mm6713e1_w
- CDC Candida auris webpage
 - https://www.cdc.gov/fungal/diseases/candidiasis/candidaauris.html
- Find your state HAI Coordinator and AR expert
 - https://www.cdc.gov/hai/state-based/index.html



Thank you!

Questions?

MollonN@Michigan.gov ipf8@cdc.gov

For more information, contact CDC 1-800-CDC-INFO (232-4636)

TTY: 1-888 -737 5248 www radc.gov



http://www.cdc.gov/longtermcare

The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the Centers for Disease Control and Prevention.



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